

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Currently Amended) A method for identifying oligonucleotide sequences suitable for the amplification of a unique sequence within a genomic region of interest, said method comprising the steps of:

executing a first process on a digital computer to identify repeat sequence-free subsequences that occur within said genomic region of interest;

executing a second process on a digital computer to compare the repeat sequence-free subsequences within said genomic region of interest to a nucleotide sequence database, whereby nucleotide sequences within said nucleotide sequence database that are at least 50% identical to said repeat sequence-free subsequences are identified;

executing a third process on a digital computer to identify, based on the repeat sequence-free subsequences, oligonucleotide sequences that are suitable for use as primers in an amplification reaction to amplify a product within at least one of said repeat sequence-free subsequences, wherein the nucleotide sequence database contains 5 or fewer sequences at least 50% identical to the repeat-free subsequence from which the oligonucleotide sequence is identified ~~for which there are 5 or fewer sequences at least 50% identical to a nucleotide sequence in said nucleotide sequence database;~~ and

outputting said oligonucleotide sequences.

2. (Original) The method of claim 1, wherein said genomic region is from a human genome.

3. (Previously Amended) The method of claim 1, wherein the third process comprises identifying oligonucleotide sequences that are suitable for use as primers in an amplification reaction to amplify a product within at least one repeat sequence-free subsequence,

wherein the database has no sequences at least 50% identical to the repeat sequence-free subsequence.

4. (Original) The method of claim 1, wherein said oligonucleotide sequences are outputted by displaying the sequences on a computer screen or on a computer printout.

5. (Original) The method of claim 1, wherein said oligonucleotide sequences are outputted by executing a fourth process on a digital computer to direct the synthesis of oligonucleotide primers comprising said oligonucleotide sequences.

6. (Original) The method of claim 5, wherein said computer directs the synthesis of said oligonucleotide primers by ordering said synthesis from an external source.

7. (Original) The method of claim 5, wherein said computer is in communication with an oligonucleotide synthesizer, and wherein said computer directs the synthesis of said oligonucleotide primers by said synthesizer.

8. (Canceled)

9. (Previously Amended) The method of claim 1, wherein the third process comprises identifying oligonucleotide sequences that are suitable for use as primers in an amplification reaction to amplify a product within at least one repeat sequence-free subsequence, wherein the database has no sequences at least 70% identical to the repeat sequence-free subsequence.

10. (Previously Amended) The method of claim 1, wherein the third process comprises identifying oligonucleotide sequences that are suitable for use as primers in an amplification reaction to amplify a product within at least one repeat sequence-free subsequence, wherein the database has no sequences at least 90% identical to the repeat sequence-free subsequence.

11. (Previously Amended) The method of claim 1, wherein said first process is executed using a software program that screens sequences for:

- i. interspersed repeats that are known to exist in mammalian genomes and;
- ii. low complexity DNA sequences.

12. (Previously Amended) The method of claim 1, wherein said second process is executed using a sequence comparison algorithm.

13. (Previously Amended) The method of claim 1, wherein said third process is executed using primer design software.

14. (Original) The method of claim 5, further comprising producing an amplification product using said oligonucleotide primers.

15. (Original) The method of claim 14, wherein said amplification product is a FISH probe.

16. (Original) The method of claim 15, wherein said FISH probe is fluorescently labeled.

17. (Original) The method of claim 14, wherein said amplification product is an array CGH target.

18. (Currently Amended) A method for identifying oligonucleotide sequences suitable for the amplification of a unique sequence within a genomic region of interest, said method comprising the steps of:

- identifying repeat sequence-free subsequences within said genomic region;
- comparing at least one repeat sequence-free subsequence within said genomic nucleotide sequence to a nucleotide sequence database to identify sequences within said database that are at least 50% identical to said repeat sequence-free subsequence;

~~for at least one of said repeat sequence free subsequences for which 5 or fewer sequences that are at least 50% identical are identified within said nucleotide sequence database,~~
selecting, based on the identified sequences, oligonucleotide sequences that are suitable for use as primers in an amplification reaction to amplify a product within said repeat sequence-free subsequence, wherein the nucleotide sequence database contains 5 or fewer sequences at least 50% identical to the repeat-free subsequence from which the oligonucleotide sequence is identified.

19. (Original) The method of claim 18, wherein said genomic region is from a human genome.

20. (Previously Amended) The method of claim 18, wherein said oligonucleotide sequences that are suitable for use as primers in an amplification reaction to amplify a product within said repeat sequence-free subsequence are selected from at least one of said repeat sequence-free subsequences that lack any sequences that are at least 50% identical to said nucleotide sequence database.

21. (Currently amended) The method of claim 18, further comprising displaying said selected oligonucleotide sequences on a computer screen or on a computer printout.

22. (Currently amended) The method of claim 18, further comprising directing the synthesis of oligonucleotide primers comprising said selected oligonucleotide sequences.

23. (Original) The method of claim 22, wherein said synthesis is directed by ordering the synthesis of said primers from an external source.

24. (Canceled)

25. (Previously Amended) The method of claim 18, comprising selecting oligonucleotide sequences that are suitable for use as primers in an amplification reaction to

amplify a product within a repeat sequence-free subsequence, wherein the database has no sequences at least 70% identical to the repeat sequence-free subsequence.

26. (Previously Amended) The method of claim 18, comprising selecting oligonucleotide sequences that are suitable for use as primers in an amplification reaction to amplify a product within a repeat sequence-free subsequence, wherein the database has no sequences at least 90% identical to the repeat sequence-free subsequence.

27. (Previously Amended) The method of claim 18, wherein the identification of repeat sequences within said genomic region is performed using a software program that screens sequences for:

- i. interspersed repeats that are known to exist in mammalian genomes and;
- ii. low complexity DNA sequences.

28. (Previously Amended) The method of claim 18, wherein the comparison of said at least one repeat sequence-free subsequence with said genome database is performed using a sequence comparison algorithm.

29. (Previously Amended) The method of claim 18, wherein said oligonucleotide sequences are selected using primer design software.

30. (Original) The method of claim 22, further comprising generating an amplification product using said oligonucleotide primers.

31. (Original) The method of claim 30, wherein said amplification product is a FISH probe.

32. (Original) The method of claim 31, wherein said FISH probe is fluorescently labeled.

33. (Original) The method of claim 30, wherein said amplification product is an array CGH target.

34. (Currently Amended) A computer program product designing and outputting oligonucleotide sequences suitable for use as primers to amplify unique sequences within a genomic region of interest, said computer program product comprising:

a storage structure having computer program code embodied therein, said computer program code comprising:

computer program code for causing a computer to identify repeat sequence-free subsequences within said nucleotide sequence;

computer program code for causing a computer to, for each subsequence of said nucleotide sequence that does not contain any of said repeat sequences, compare said subsequence against a nucleotide sequence database to identify nucleotide sequences within said database that are at least 50% identical to said subsequence;

computer program code for causing a computer to, ~~for at least one of said subsequences for which 5 or fewer sequences that are at least 50% identical are found in said database,~~ identify oligonucleotide sequences suitable for use as primers in an amplification reaction to amplify a product within said subsequence, wherein the nucleotide sequence database contains 5 or fewer sequences at least 50% identical to the repeat-free subsequence from which the oligonucleotide sequence is identified; and

computer program code for outputting said oligonucleotide sequences.

35. (Previously Amended) The method of claim 34, wherein the computer program product comprises code for causing a computer to identify oligonucleotide sequences that are suitable for use as primers in an amplification reaction to amplify a product within said subsequence, wherein the database has no sequences at least 50% identical to the subsequence.

36. (Canceled)

37. (Previously Amended) The method of claim 34, wherein the computer program product comprises code for causing the computer to select oligonucleotide sequences that are suitable for use as primers in an amplification reaction to amplify a product within a

repeat sequence-free subsequence, wherein the database has no sequences at least 70% identical to the repeat sequence-free subsequence.

38. (Previously Amended) The method of claim 34, wherein the computer program product comprises code for causing the computer to select oligonucleotide sequences that are suitable for use as primers in an amplification reaction to amplify a product within a repeat sequence-free subsequence, wherein the database has no sequences at least 90% identical to the repeat sequence-free subsequence.

39. (Canceled)

40. (Previously Added) The method of claim 1, wherein the repeat-free subsequences are each at least 100 bp long.

41. (Previously Added) The method of claim 18, wherein the repeat-free subsequences are each at least 100 bp long.

42. (Previously Added) The computer program of claim 34, wherein each nucleotide sequence that does not contain any of the repeat sequences is at least 100 bp long.

43. (Currently Amended) A method for identifying oligonucleotide sequences suitable for the amplification of a unique sequence within a genomic region of interest, said method comprising the steps of:

executing a first process on a digital computer to identify repeat sequence-free subsequences that occur within said genomic region of interest;

executing a second process on a digital computer to compare repeat sequence-free subsequences within said genomic region of interest to a nucleotide sequence database, whereby at least one repeat sequence-free subsequence that is at least 50% identical to a nucleotide sequence within said nucleotide sequence database is discarded;

executing a third process on a digital computer to identify, based on the repeat sequence-free subsequences, oligonucleotide sequences that are suitable for use as primers in an

amplification reaction to amplify a product within at least one repeat sequence-free subsequences remaining after executing said second process; and
outputting said oligonucleotide sequences.

44. (Previously Added) The method of claim 43, wherein said genomic region is from a human genome.

45. (Previously Added) The method of claim 43, wherein said oligonucleotide sequences are outputted by displaying the sequences on a computer screen or on a computer printout.

46. (Previously Added) The method of claim 43, wherein said oligonucleotide sequences are outputted by executing a fourth process on a digital computer to direct the synthesis of oligonucleotide primers comprising said oligonucleotide sequences.

47. (Previously Added) The method of claim 43, wherein said computer directs the synthesis of said oligonucleotide primers by ordering said synthesis from an external source.

48. (Previously Added) The method of claim 43, wherein said computer is in communication with an oligonucleotide synthesizer, and wherein said computer directs the synthesis of said oligonucleotide primers by said synthesizer.

49. (Previously Added) The method of claim 43, wherein all repeat sequence-free subsequences that are at least 70% identical to a nucleotide sequence within said nucleotide sequence database are discarded.

50. (Previously Amended) The method of claim 43, wherein all repeat sequence-free subsequences that are at least 90% identical to a nucleotide sequence within said nucleotide sequence database are discarded.

51. (Previously Added) The method of claim 43, wherein said first process is executed using a software program that screens sequences for:

- i. interspersed repeats that are known to exist in mammalian genomes and;
- ii. low complexity DNA sequences.

52. (Previously Added) The method of claim 43, wherein said second process is executed using a sequence comparison algorithm.

53. (Previously Amended) The method of claim 43, wherein said third process is executed using primer design software.

54. (Previously Added) The method of claim 43, further comprising producing an amplification product using said oligonucleotide primers.

55. (Previously Added) The method of claim 43, wherein said amplification product is a FISH probe.

56. (Previously Added) The method of claim 43, wherein said FISH probe is fluorescently labeled.

57. (Previously Added) The method of claim 43, wherein said amplification product is an array CGH target.

58. (Previously Added) The method of claim 43, wherein the repeat-free subsequences are each at least 100 bp long.

59. (Previously Added) The method of claim 43, wherein all repeat sequence-free subsequences that are at least 90% identical to a nucleotide sequence within said nucleotide sequence database are discarded.

60. (Currently Amended) A method for identifying oligonucleotide sequences suitable for the amplification of a unique sequence within a genomic region of interest, said method comprising the steps of:

- (1) identifying repeat sequence-free subsequences repeat sequences within said genomic region;
- (2) comparing repeat sequence-free subsequences within said genomic region of interest to a nucleotide sequence database, whereby at least one repeat sequence-free subsequences that is at least 50% identical to a nucleotide sequence within said nucleotide sequence database is discarded;
- (3) identifying oligonucleotide sequences, based on the repeat sequence-free subsequences, that are suitable for use as primers in an amplification reaction to amplify a product within at least one repeat sequence-free subsequences remaining after step (2).

61. (Currently Amended) A computer program product designing and outputting oligonucleotide sequences suitable for use as primers to amplify unique sequences within a genomic region of interest, said computer program product comprising a storage structure having computer program code embodied therein, said computer program code comprising the elements:

- (1) computer program code for causing a computer to identify repeat sequence-free subsequences within said nucleotide sequence;
- (2) computer program code for causing a computer to compare repeat sequence-free subsequences within said genomic region of interest to a nucleotide sequence database, whereby at least one repeat sequence-free subsequences that is at least 50% % identical to a nucleotide sequence within said nucleotide sequence database is discarded;
- (3) computer program code for causing a computer to identify, based on the repeat sequence-free subsequences, oligonucleotide sequences that are suitable for use as primers in an amplification reaction to amplify a product within at least one repeat sequence-free subsequences remaining after executing said computer program code in element (2); and
- (4) computer program code for outputting said oligonucleotide sequences.